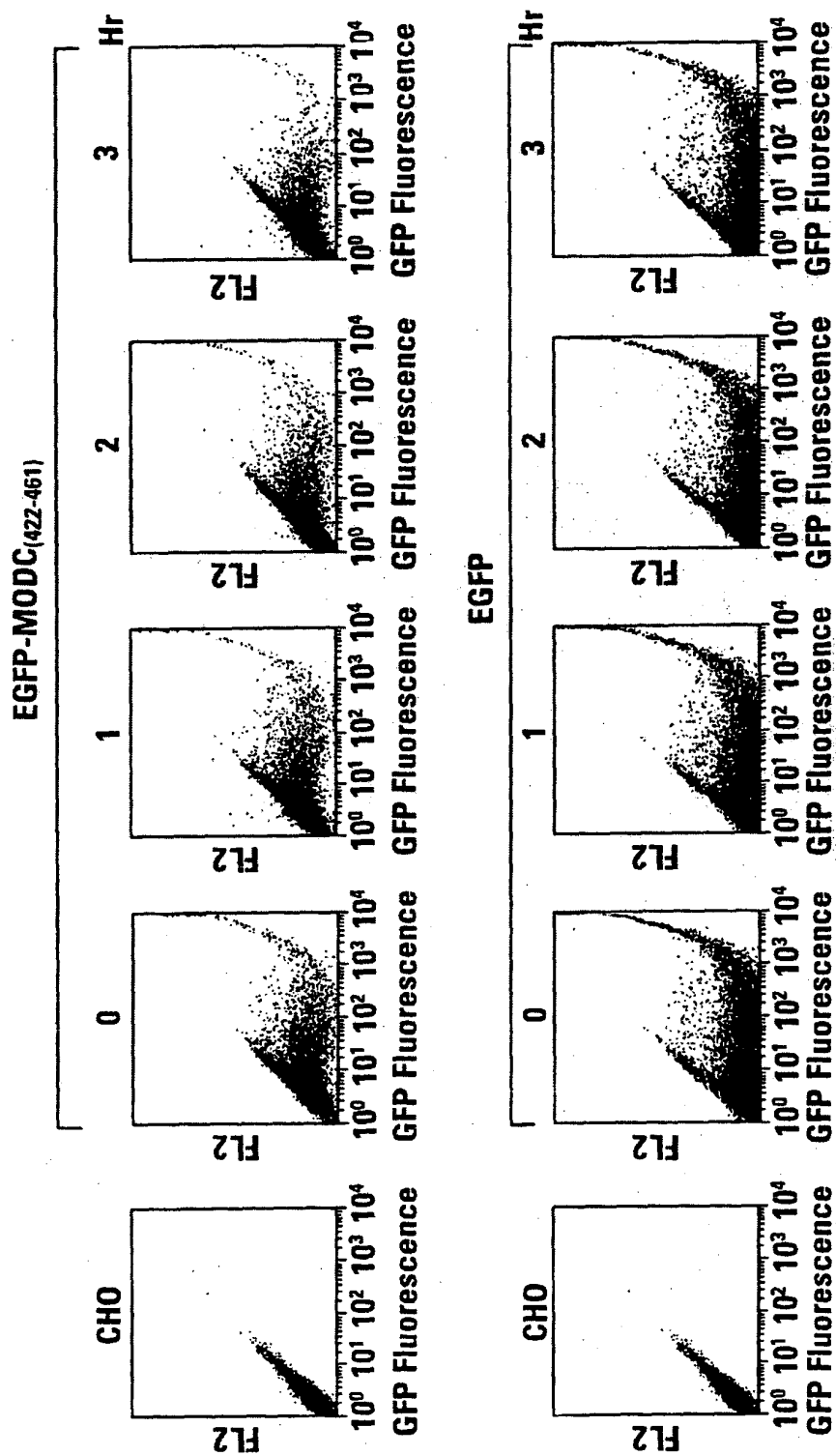


FIG. 1



FIG. 2



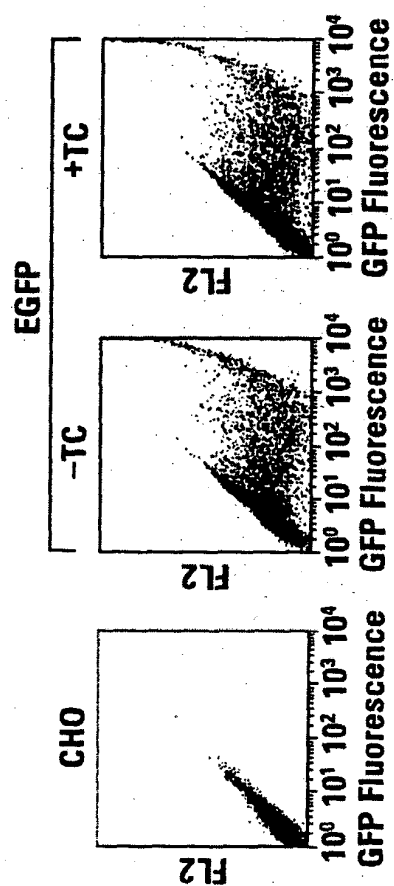


FIG. 3 cont.

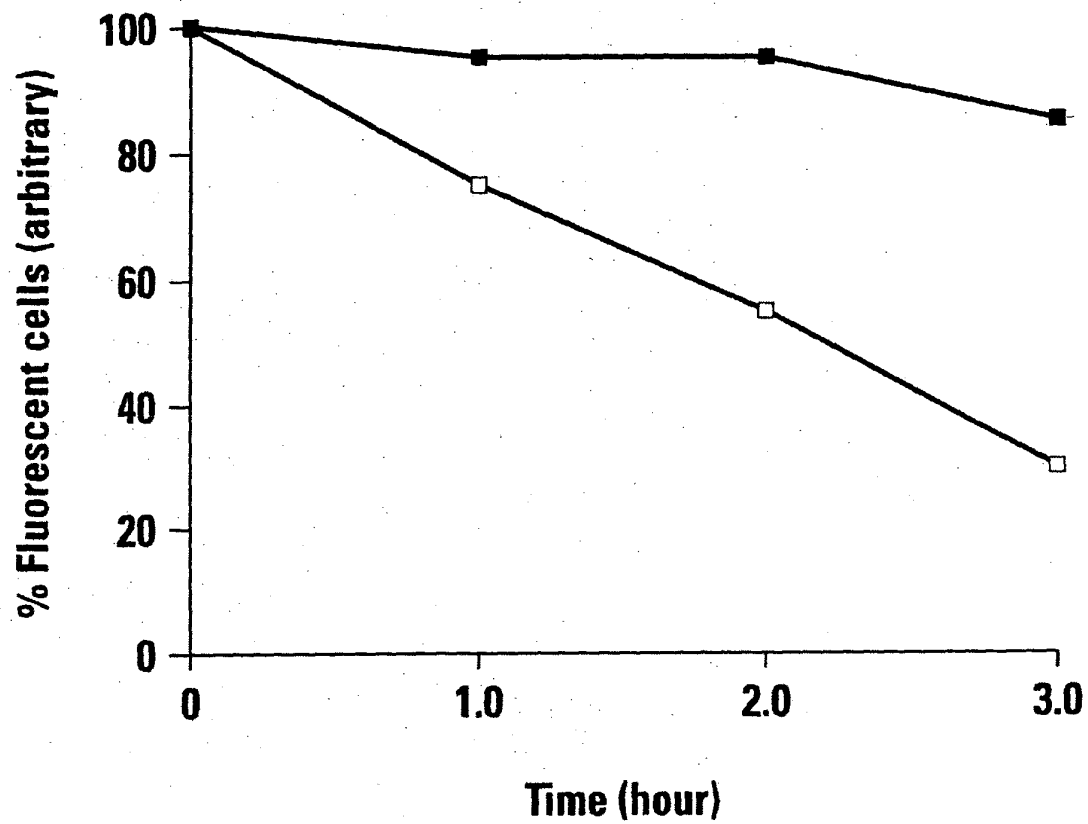


FIG. 4

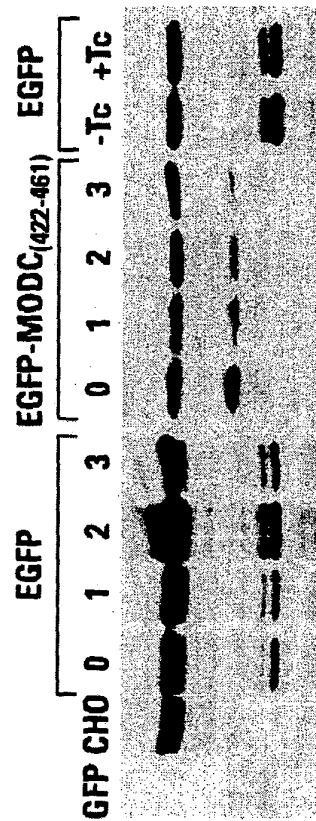
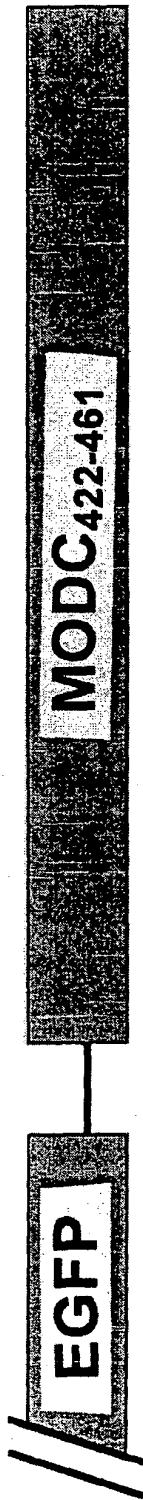


FIG. 5



PEST Sequence

	423	450
EGFP-MODC <sub>422-461</sub>	--HGFPPEVEEQDDGTLFMSCAQESGMDRH--	(SEQ ID NO. 3)
P426A/P427A	---AA---	
P438A	-----A-----	
E428A/E430A/E431A	-----A-AA-----	
E44A	-----A-----	
S440A	-----A-----	
S445A	-----A-----	
T436A	-----A-----	
D433A/D434A	-----AA-----	
D448A	-----A-----	
H423A	---A-----	
R449A/H450A	-----AA-----	

FIG. 6

## FIG. 7

**Table 1. FACS analysis of EGFP, EGFP-MODC<sup>422-461</sup>, and mutations in transfected CHO K1 Tet-off cells.**

Constructs	0h	(initial)	2h	4h
EGFP	100%	(63.6)	107%	92%
EGFP-MODC <sup>422-461</sup>	100%	(12.6)	52%	29%
P426A/P427A	100%	(11.5)	39%	11%
P438A	100%	(34.1)	79%	60%
E428A/E430A/E431A	100%	(17.3)	20%	15%
E444A	100%	(12.6)	69%	65%
S440A	100%	(21.6)	78%	66%
S445A	100%	(23.5)	29%	20%
T436A	100%	(46.9)	70%	47%
D433A/D434A	100%	(11.31)	22%	6%
D448A	100%	(32.6)	30%	15%
H423A	100%	(12.2)	50%	25%
R449A/H450A	100%	(27.9)	93%	86%

Transfection was performed in CHO/tTA cells using the procedure as described in Methods. After 24 hours, cells were treated with CHx for 0, 2 and 4 hours, and analyzed for fluorescence intensity by FACS Caliber (Becton Dickinson). The fluorescent cells at each time point are represented as a percentage of initial.



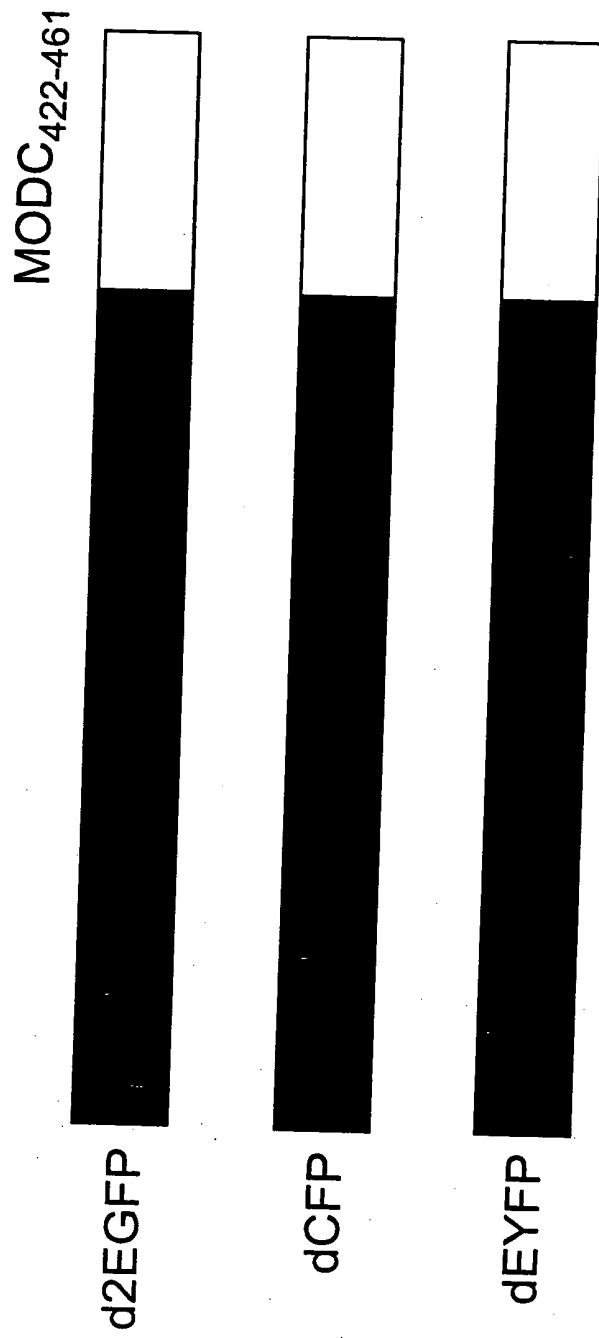


FIG. 8

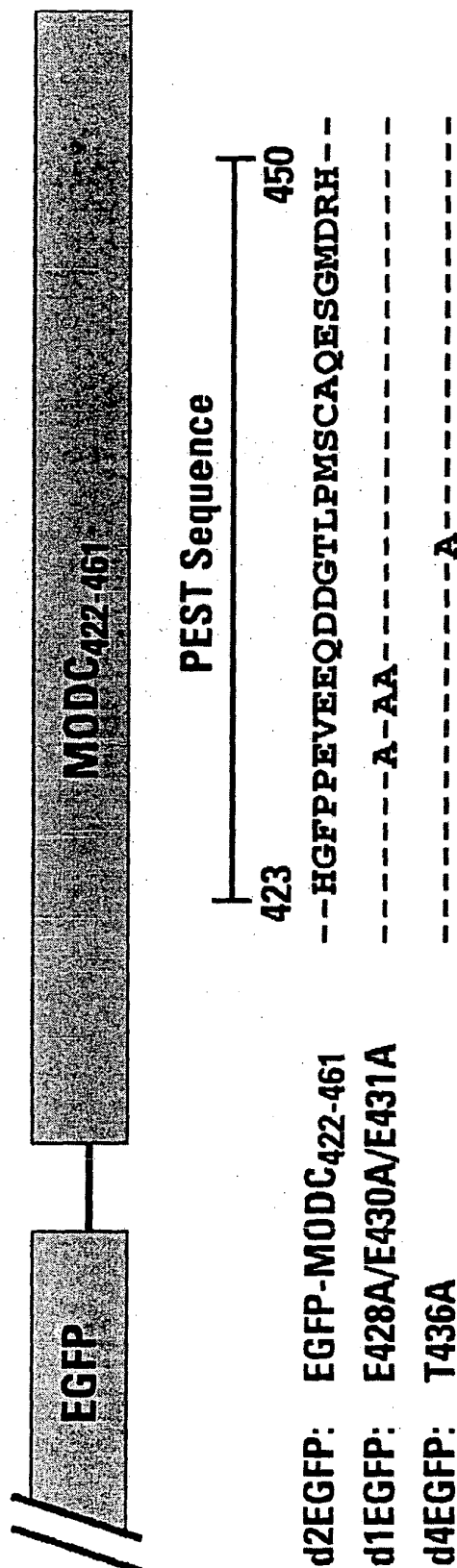


FIG. 9

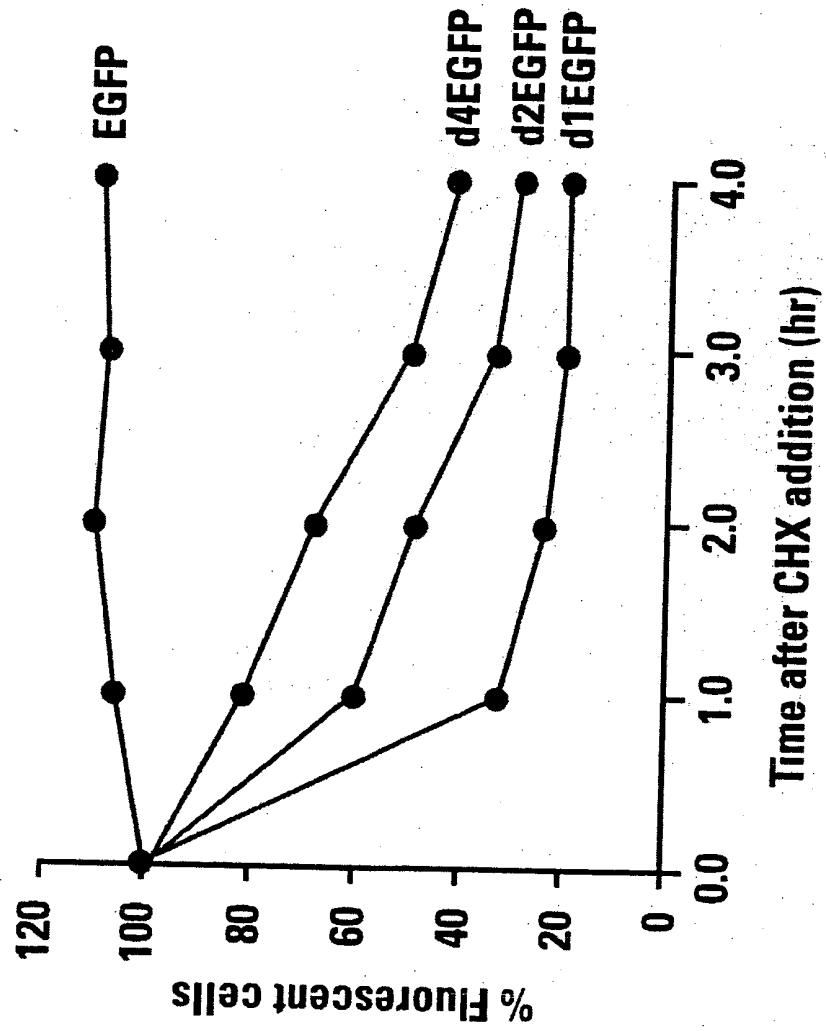


FIG. 10

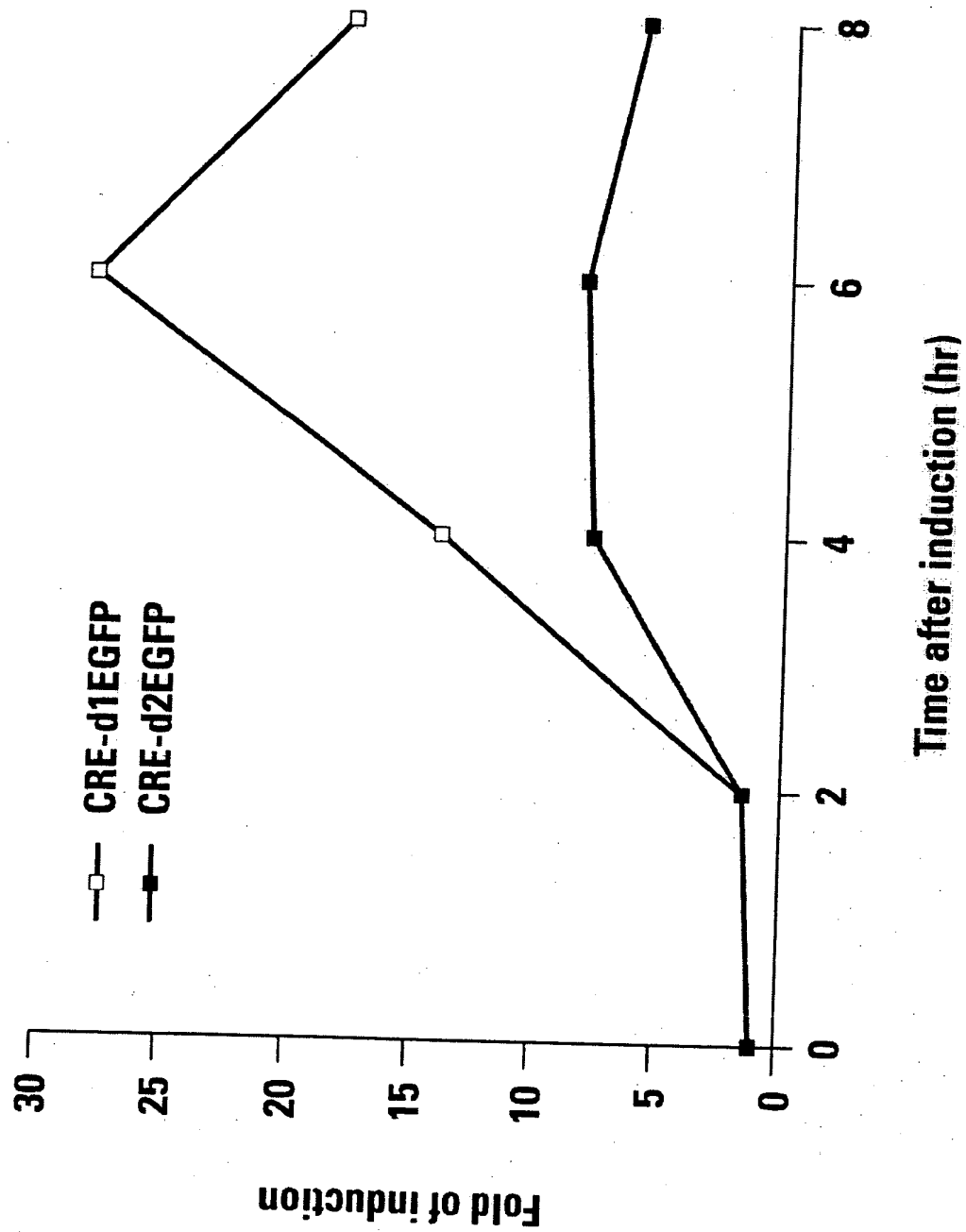


FIG. 11